

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE
REQUEST FOR FILING NATIONAL PATENT APPLICATION

Under 35 USC 111(a) and Rule 53(b)

PATENT APPLICATION

Asst. Commissioner of Patents
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WITH SIGNED DECLARATION

JC685 U.S. PTO



09/15/00

NONPROVISIONAL
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NON PCT NAT PHASE

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09/662937



Sir:

Herewith is the PATENT APPLICATION of
Inventor(s): KITAGAWA et al.

Title BRIGHT SURFACE STRUCTURE AND A MANUFACTURING
METHOD THEREOF

Atty. Dkt.: PM 273286
M#

US990172
Client Ref

Date: September 15, 2000

including:

1. Specification: 36 pages (only spec. and claims) 2. ☐ Specification in non-English language
3. Declaration ☒ Original ☐ Facsimile/Copy ☒ Abstract 1 page(s); 38 numbered claims
4. ☒ Drawings: 1 sheet(s) ☐ informal; ☒ formal of size: ☒ A4 ☐ 11"
5. ☐ See top first page re prior Provisional, National or International application(s). ("X" box only if info is there and do not
complete corresponding item 5 or 6). (Prior M# _____ SN _____)
6. AMEND the specification please by inserting before the first line: -- This is a ☐ Continuation-in-Part
☐ Divisional ☐ Continuation ☐ Substitute Application (MPEP 201.09) of:
6(a) ☐ National Appln. No. _____ / _____ filed _____ (M# _____)
6(b) ☐ International Appln. No. _____ filed _____
7. ☐ AMEND the specification by inserting before the first line: -- This application claims the benefit of U.S.
Provisional Application No. 60/ _____, filed _____
8. ☒ Attached is an assignment and cover sheet. Please return the recorded assignment to the undersigned.
9. ☐ Prior application is assigned to _____

by Assignment recorded _____ Reel _____ Frame _____

10. FOREIGN priority is claimed under 35 USC 119(a)-(d)/365(b) based on filing in Japan

11. _____ (country)

Application No.	Filing Date	Application No.	Filing Date
(1) 11-264172	September 17, 1999	(2) 11-270072	September 24, 1999
(3)		(4)	
(5)		(6)	
(7)		(8)	
(9)		(10)	

12. _____ (No.) Certified copy (copies): ☐ attached; ☐ previously filed (date) _____
in U.S. Application No. _____ / _____ filed on _____

13. ☐ Attached: _____ (No.) Verified Statement(s) establishing "small entity" status under Rules 9 & 27.

14. **DOMESTIC/INTERNATIONAL** priority is claimed under 35 USC 119(e)/120/365(c) based on the following provisional, nonprovisional and/or PCT international application(s):

Application No.	Filing Date	Application No.	Filing Date
(1)		(4)	
(2)		(5)	
(3)		(6)	

15. ☐ This application is being filed under Rule 53(b)(2) since an inventor is named in the enclosed Declaration who was not named in the prior application.

16. ☒ Attached: Form PTO-1449 and a copy of the listed documents. See this application, e.g., page 2, for comments.

17. ☐ Preliminary Amendment:

THE FOLLOWING FILING FEE IS BASED ON CLAIMS AS FILED LESS ANY ABOVE CANCELLED

				Large/Small Entity		Fee Code
18. Basic Filing Fee				\$690/\$345	\$690	101/201
19. Total Effective Claims	70	minus 20 =	*50	x \$18/\$9 =	+ 900	103/203
20. Independent Claims	4	minus 3 =	*1	x \$78/\$39 =	+ 78	102/202
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21. If any proper multiple dependent claim (ignore improper) is present, add (Leave this line blank if this is a reissue application)				+ \$260/\$130	+ 260	104/204
22.	TOTAL FILING FEE ENCLOSED =				\$1928	
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26.	TOTAL FEE ENCLOSED =				\$1968	

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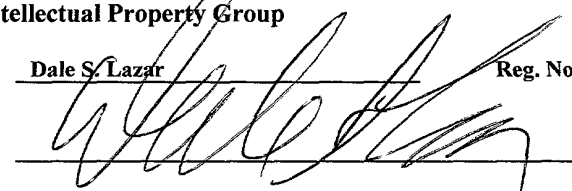
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APPLICATION UNDER UNITED STATES PATENT LAWS

Atty. Dkt. No. PM 273286
(M#)

Invention: BRIGHT SURFACE STRUCTURE AND A MANUFACTURING METHOD THEREOF

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- ☐ Provisional Application
- ☒ Regular Utility Application
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SPECIFICATION

BRIGHT SURFACE STRUCTURE AND A MANUFACTURING METHOD THEREOF

This application is based on and claims priority from Japanese Patent
Application Nos. H11-264172 filed on September 17, 1999 and H11-270072 filed
on September 24, 1999, the contents of which are incorporated herein by
reference.

BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention relates to a bright surface structure formed on a
member made from metal or resin and having a good ductility and corrosion
resistance, and a method for manufacturing thereof.

2. Description of Related Art

To brighten a surface of a metal or a resin member, conventionally an
electro plating, such as an electro chromium plating is used. By this method, a
high reflectivity, a gloss of a high degree having an appearance of deepness, a
good wearing resistance and a good corrosion resistance are obtained. This
surface brightening treatment is applied to a metal member such as an aluminum
wheel of an automobile, and a resin member such as a front grille, a garnish and
an emblem of an automobile for lightening the member.

Since the electro plating needs an environmental countermeasure, recently
various methods replacing the electro plating have been investigated. As one
substituting method, with the aluminum wheel of an automobile, a surface of the
wheel is ground to a smooth surface. However, the grinding is not only

accompanied by an increase in manufacturing cost but also inapplicable to an inside surface of ventilation holes of the wheel.

Japanese Patent Publications H04-131232 and H10-130822 propose the following improving methods replacing the grinding method:

5 (1) Japanese Patent Publication H04-131232

After a metal member is surface-treated, the member is coated with an under coat, then a thin chromium film formed by sputtering, and then a top coat. A surface of the thin chromium film operates as a reflective surface. In the surface treatment, the surface of the member is shot-blasted and coated with a clear resin by powder coating. As the under coat, a clear resin film is formed. The chromium film is formed using chromium. The top coat is a clear coat that protects bright surface of the chromium film.

However, the method described in Japanese Patent Publication H04-131232 presents a problem of the thin chromium film cracking, though thin chromium film presents an external appearance similar to electro chromium plating. Because the residual stress of the chromium film is large, cracking is caused during sputtering and after the top coat is coated, or after the top coat is coated on the chromium film and dried. When the thickness of the chromium film is equal to or greater than $0.04\ \mu\text{m}$, the crack initiation is severe. The residual stress is so significant that when a stress generated in the top coat during abrasion test and weather resistance test is transmitted to the chromium film, cracking easily occurs in the chromium film. Further, when a solvent penetrates the chromium film during coating and/or drying the top coat, cracking easily occurs in the chromium film.

25 If the chromium film is reduced in thickness in order to suppress

generation of the cracks in the chromium film, the external appearance similar to electro chromium plating cannot be obtained any longer because an under wheel surface is seen through the thin chromium coat and the brightness of chromium film is reduced.

5 (2) Japanese Patent Publication H10-130822

After a powder coating is applied to an aluminum forging part, an under coat, then a thin aluminum film formed by sputtering, and then a top coat are formed. A surface of the thin aluminum film operates as a reflective surface.

However, the method described in Japanese Patent Publication
10 H10-130822 presents a problem that corrosion resistance is relatively low, though no cracks occur even if the aluminum film has a thickness of about $0.08 \mu\text{m}$, since aluminum film has a high ductility. Though the top resin coat formed on the aluminum film protects the aluminum film, once the top coat is damaged during lifetime of an automobile, corrosion propagates from the damaged top coat.
15 When the aluminum film is removed due to the corrosion, the brightness is gone.

SUMMARY OF THE INVENTION

The present invention provides for a bright surface structure formed on a metal or resin member having an external surface similar to an electro chromium
20 plating, and a good ductility and corrosion resistance, and a method for manufacturing thereof.

BRIEF DESCRIPTION OF THE DRAWINGS

The above discussed and other objects, features, and advantages of the

present invention will become more apparent and will be more readily appreciated from the following detailed description of the preferred embodiments of the present invention in conjunction with the accompanying drawings, in which:

FIG. 1 is a cross-sectional view of a bright surface structure according to the first embodiment of the present invention and manufactured by a method according to the fourth embodiment of the present invention;

FIG. 2 is a cross-sectional view of a bright surface structure according to the second embodiment of the present invention and manufactured by a method according to the fifth embodiment of the present invention; and

FIG. 3 is a cross-sectional view of a bright surface structure according to the third embodiment of the present invention and manufactured by a method according to the sixth embodiment of the present invention.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

FIG.1 illustrates a bright surface structure according to the first embodiment of the present invention and obtained by a manufacturing method for a bright surface structure according to the fourth embodiment of the present invention. FIG.2 illustrates a bright surface structure according to the second embodiment of the present invention and obtained by a manufacturing method for a bright surface structure according to the fifth embodiment of the present invention. FIG.3 illustrates a bright surface structure according to the third embodiment of the present invention and obtained by a manufacturing method for a bright surface structure according to the sixth embodiment of the present invention.

Portions common or similar to each other throughout all of the

embodiments of the present invention are denoted with the same reference numerals throughout all of the embodiments of the present invention.

As illustrated in FIG. 1, a bright surface structure formed on a member 4 made from metal or resin according to the first embodiment of the present invention includes (a) a resin film 1 coated on the member 4, and (b) a thin metal film 2 formed on the resin film 1. The thin metal film 2 is made from material selected from the group consisting of stainless steel, titanium alloy and nickel alloy and has a smooth surface.

As illustrated in FIG. 2, a bright surface structure formed on a member 4 made from metal or resin according to the second embodiment of the present invention further includes (c) a clear (transparent) protective film 3 coated on the thin metal film 2, in addition to the bright surface structure according to the first embodiment of the present invention.

As illustrated in FIG. 3, a bright surface structure formed on a member 4 made from metal or resin according to the third embodiment of the present invention includes (a) a resin film 1 coated on the member 4, (b') a thin metal film 2' formed on the resin film 1, and (c') a clear (transparent) colored protective film 3' coated on the thin metal film. The thin metal film 2' is made from material selected from the group consisting of stainless steel, titanium alloy, nickel alloy, aluminum, titanium and chromium and has a smooth surface.

As illustrated in FIG. 1, a method for manufacturing a bright surface structure formed on a member 4 made from metal or resin according to the fourth embodiment of the present invention includes (a) coating a resin film 1 on the member 4 such that the resin film 1 has a smooth surface, (b) forming a thin metal film 2 on the resin film 1. The thin metal film 2 is made from material

selected from the group consisting of stainless steel, titanium alloy and nickel alloy and has a smooth surface.

As illustrated in FIG. 2, a method according to the fifth embodiment of the present invention further includes (c) coating a clear protective film 3 on the metal film 2, in addition to the method according to the fourth embodiment of the present invention.

As illustrated in FIG. 3, a method for manufacturing a bright surface structure formed on a member 4 made from metal or resin according to the sixth embodiment of the present invention includes (a) coating a resin film 1 on the member 4 such that the resin film 1 has a smooth surface, (b') forming a thin metal film 2' on the resin film 1, and (c') coating a clear colored protective film 3' on the thin metal film 2'. The thin metal film 2' is made from material selected from the group consisting of stainless steel, titanium alloy and nickel alloy, aluminum, titanium and chromium and has a smooth surface.

In any of the first through the sixth embodiments of the present invention, the member 4 may be a metal member, such as an aluminum wheel of an automobile, or a resin member, such as a front grille, a garnish and an emblem of an automobile.

In the preferred embodiments, the stainless steel is austenitic stainless steel; the titanium alloy contains 20–80% by weight of titanium and 20–80% by weight of aluminum; the nickel alloy contains 30–80% by weight of nickel and 15–25% by weight of chromium.

Preferably, when the thin metal film 2 or 2' is made from stainless steel or titanium alloy, the thin metal film 2, 2' has a thickness of 0.03–1.0 μm , and when the thin metal film 2 or 2' is made from a nickel alloy, the thin metal film 2 or 2'

has a thickness of 0.03–0.5 μ m.

Preferably, the clear protective film 3 has a thickness of 5–20 μ m.

Preferably, the clear colored protective film 3' has a thickness of 20–40 μ m.

5 Preferably, the clear colored protective film 3' is made from a clear resin comprising a pigment or a dye. The clear resin is any one of acrylic resin-based, urethan-based and epoxy-based resins. The pigment is any one of carbon-based, lead chromate-based, iron(II) ferrocyanide-based, cobalt-based, and chromium oxide-based pigments. The pigment is also any one of thren-based, quinacrine
10 staining-based, isoindolinone-based or metal complex pigment. The dye is any one of acid dyes, mordant dyes, basic dyes, disperse dyes, edible dyes, direct dyes and sulphur dyes.

Preferably, the resin film 1 is formed by powder coating.

Preferably, the thin metal film 2, 2' is formed by cathode-arc-type ion
15 plating or sputtering.

Each of the films 1, 2, 2', 3 and 3' in the first through the sixth embodiments of the present invention will be explained in more detail.

(a) Resin Film 1

The resin film 1 coated on the metal or resin member 4 is required to have
20 a good adhesion to the member 4, to have a smooth surface so that the thin metal film 2, 2' formed on the resin film 1 has a smooth surface and has a high reflectivity, and to have a good adhesion with the thin metal film 2, 2'. Any one of polyester-based, epoxy-based, acrylic resin-based, and urethan-based resins can be used as material for the resin film 1. Any color may be used. As long as
25 the above requirement is satisfied, the resin film 1 may be of any thickness.

(b) Thin Metal Film 2 and (b') Thin Metal Film 2'

A material for the thin metal film 2 is selected from stainless steel, titanium alloy and nickel alloy. The thin metal film 2 made from any one of these materials has an external appearance similar to that of electro chromium plating and has a good ductility (crack resistance) and a good wear resistance.

As the stainless steel, austenitic stainless steel is preferable. Martensitic stainless steel and ferritic stainless steel is inferior to austenitic stainless steel in an external appearance, ductility and wear resistance. Therefore, a thickness of the thin metal film 2, 2' is determined taking into account the kind of stainless steel used for the thin metal film.

It is preferable that the titanium alloy for the thin metal film 2, 2' contains 20–80% by weight of titanium. In case the alloy contains less than 20% by weight of titanium, the wear resistance of the thin metal film 2, 2' will decrease though the brightness is maintained, so that during salt water spray test, the metal film will be melted. In case the alloy contains more than 80% by weight of titanium, the brightness of the thin metal film 2, 2' is suppressed and the color of the thin metal film 2, 2' approaches that of chromium, and during warm water immersion test, an oxidized film is generated at the surface to produce an interference color. The titanium alloy containing 20–80% by weight of aluminum is more preferable. The contained aluminum makes higher ductility and reflectivity of the titanium alloy.

It is preferable that the nickel alloy for the thin metal film 2, 2' contains 30–80% by weight of nickel. In case the alloy contains less than 30% by weight of nickel, the color of the thin metal film 2, 2' approaches the chromium color, suppressed in brightness and becomes harder to decrease in crack resistance. In

case the alloy contained more than 80% by weight of nickel, the thin metal film 2, 2' has excessive brightness. A nickel-chromium alloy containing about 30% by weight of nickel and 15-25% by weight of chromium is more preferable.

With respect to a thickness of the thin metal film 2, 2', in case of the stainless steel film and the titanium alloy film, a thickness of 0.03-1.0 μm is preferable, and in case of nickel alloy film, 0.03-0.5 μm is preferable. In case a thickness is less than 0.03 μm , the under resin film 1 can be seen through the metal film and a high reflectivity cannot be achieved. In case the thickness is more than the above respective upper limits, cracks are generated due to the residual stress in the film, and further, a time period for forming the thin metal film becomes long and the productivity decreases.

(c) Clear Protective Film 3

In order to increase the wear resistance and the corrosion resistance and to obtain an external appearance similar to that of electro chromium plating, a clear (transparent) protective film 3 is formed on the thin metal film 2. The clear protective film 3 is required to have a good adhesion to the thin metal film 2 and a good weather resistance so as to maintain the transparency for prolonged time periods. A coating paint used for forming the clear protective film 3 is any one of acrylic resin-based, urethan-based and epoxy-based resins. A thickness of the clear protective film 3 is conventionally 20-40 μm , but in the present invention, since the thin metal film 2 has a high corrosion resistance, a thickness of the clear protective film 3 can be 5-20 μm and obtain satisfactory wear resistance. Further, since the entire surface of the thin metal film 2 is not necessarily coated with the clear protective film 3, the brightening treatment cost can be decreased.

(c') Clear Colored Protective Film 3'

In order to increase the wear resistance and the corrosion resistance and to obtain an external appearance similar to that of electro chromium plating, a clear (transparent) colored protective film 3' is formed on the thin metal film 2. The clear colored protective film 3' is required to have a good adhesion to the thin metal film 2 and a good weather resistance so as to maintain the transparency for prolonged time periods. A coating paint used for forming the clear colored protective film 3' is any one of acrylic resin-based, urethan-based and epoxy-based resins.

The (clear resin) paint comprises a pigment or a dye to give a good external appearance to the member 4.

There are two kinds of pigments: an inorganic pigment and an organic pigment. The inorganic pigment includes a natural pigment and an artificial pigment, and are stable in color. A carbon-base offers black, a lead chromate-base offers yellow, an iron(II) ferrocyanide-base offers blue, a cobalt-base offers blue, and a chromium oxide-base offers green. The organic pigment includes a thren-based (which offers red, blue, orange and purple), a quinacrine staining-based (which offers red, magenta, purple and yellow), an isoindolinone-based (which offers yellow and red), and metal complex pigments (which offers yellow and green).

The dyes are classified into an acid dye, a mordant dye, a basic dye, a disperse dye, an edible dye, a direct dye and a sulphur dye and can offer yellow, orange, red, purple, blue, green, brown and black. The dyes are also classified by a chemical structure into nitro, azo, diphenyl, triphenylmethan, xanthene, azine, indigo, and anthraquinone.

The organic pigment and the dyes are especially suitable for achieving the

high reflectivity of the under thin metal film 2'. The diameter of a pigment or dye particle is preferably 0.1–30 μ m.

The thickness of the clear colored protective film 3' is 20–40 μ m.

Each step of the method according to the fourth through the sixth
5 embodiments of the present invention will be explained in more detail.

(a) Formation of the Resin Film 1

A roughness of the surface of the metal or resin member 4 varies from a mirror surface to a pear-skin. The surface is cleaned and degreased, and in some cases is conversion-treated to form an oxidized film at the surface, and then the
10 surface is coated with the resin film 1. Using a solvent based paint where a resin paint is dissolved in a solvent, a water based paint or powder paint, the resin film 1 is formed, by immersing the member 4 in the paint, injecting the paint to the member 4, or powder-coating. In the case where the member 4 has a surface with a roughness of 100–200 μ m, as in case of a cast, the powder coating is
15 preferable. In the powder coating, electrostatic coating is conducted and the adhering paint is baked at above 100 °C, so that even the surface with a large roughness is smoothed. The powder coated surface may be further sprayed by a solvent base paint to get a further smoothed surface.

(b) Formation of the Thin Metal Film 2 and Formation of the Thin Metal Film 2'

20 A cathode arc-type ion plating or sputtering is preferable for forming the thin metal film 2, 2'. The cathode arc-type ion plating is a method where by generating an arc discharge in a vacuum, a target is locally evaporated. The sputtering is a method where an argon ion collides with a target in a vacuum to eject out an atom of the target. Therefore, the same metal as the target forms the
25 thin metal film 2, 2'.

(c) Formation of the Clear Protective Film 3

To form the clear protective film 3, a solvent based paint of any one of acrylic-based, urethan-based and epoxy-based resins is injected by air onto the thin metal film 2.

5 (c') Formation of the Clear Colored Protective Film 3'

To form the clear colored protective film 3', a pigment of dye of a desired color is selected and the selected pigment or dye is added to and mixed with the solvent based paint of any one of acrylic-based, urethan-based and epoxy-based resins. Then, the paint with the pigment or dye is injected by air onto the thin
10 metal film 2'.

Tests were conducted to confirm that the bright surface structures according to the present invention have the desired properties. Among the following test examples, Examples 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24 and 26 correspond to the first and the fourth embodiments of the present invention;
15 Examples 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23 and 25 correspond to the second and the fifth embodiments of the present invention; and Examples 27 - 32 correspond to the third and the sixth embodiments of the present invention.

EXAMPLE 1

An aluminum alloy cast AC4C (Al-Si-Mg based) was used as a base
20 material (metal member 4). This material was cast, having surface roughness of 100-200 μ m.

Then, a chemical conversion coating was formed at the surface of the base material to increase a corrosion resistance. Then, an electrostatic epoxy powder coating was performed using the electric voltage condition of 40-60 kV, and the

resulting cast was baked at 170 °C for 60 minutes to form a resin film 1 of a thickness of about 120 μm. Further, to obtain a smoother surface, a coating of a solvent based acrylic resin was formed to a thickness of about 25 μm and was baked at 145°C for 60 minutes.

5 Then, a thin metal film 2 was formed by DC-magnetron-sputtering using a target of SUS304. The target electric current was 3A, and the vacuum pressure was 2.5×10^{-3} Torr. By coating for 30 seconds, the thin metal film 2 of a thickness of about 0.06 μm was obtained.

10 Then, a clear paint of acrylic resin was coated by spraying on the thin metal film 2 to a thickness of 20 μm and was baked at 80 °C for 60 minutes to obtain a clear protective film 3.

A visual test (visual inspection), a warm water immersion test and a corrosion resistance test were conducted.

15 According to the visual test, the thin metal film of stainless steel did not exhibit a crack and had an external appearance similar to that of electro chromium plating.

20 In the warm water immersion test, the test piece was immersed in warm water at 60°C for 72 hours. As a result of the test, there was no change in the external appearance. Further, a tape was attached to the surface and then detached from the surface. As a result of the tape test, there was no separation of the clear protective film from the thin metal film.

 In the corrosion resistance test, the clear protective film was cross-cut and was subjected to a salt water spray test for 1000 hours. As a result of the test, neither corrosion of the thin stainless steel film from the cross-cut portion nor

breakage of the thin stainless steel film was observed.

EXAMPLE 2

5 The same test piece as that of Example 1 was prepared, except that the clear protective film was not formed after formation of the thin stainless steel film.

Then, the same corrosion resistance test as that of Example 1 was conducted. As a result of the test, neither corrosion of the thin metal film from the cross-cut portion nor breakage of the thin metal film was observed.

EXAMPLE 3

10 The same test piece as that of Example 1 was prepared, except that in formation of the thin metal film a target of SUS316L was used.

Then, the same tests as those of Example 1 were conducted. The visual test, the warm water immersion test and the corrosion resistance test produced the same results as those for Example 1.

15 EXAMPLE 4

The same test piece as that of Example 3 was prepared, except that the clear protective film was not formed after formation of the thin metal film of stainless steel.

20 Then, the same tests as those of Example 1 were conducted. The visual test, the warm water immersion test and the corrosion resistance test produced the same results as those for Example 1.

EXAMPLE 5

25 The same test piece as that of Example 1 was prepared, except that the preparation conditions were changed as follows. In formation of the thin metal film, a cathode arc-type ion plating apparatus was used and the thin metal film of

the thickness of about $0.8\mu\text{m}$ was formed. The argon pressure was 30 mTorr, the cathode electric current was 60A, and the coating period of time was 15 minutes.

The temperature of the base material was controlled so that the thin metal film formation temperature did not exceed the baking temperature of the under resin film.

Then, the same tests as those of Example 1 were conducted. The visual test, the warm water immersion test and the corrosion resistance test produced the same results as those for Example 1.

EXAMPLE 6

The same test piece as that of Example 5 was prepared, except that the clear protective film was not formed after formation of the thin stainless steel film.

Then, the same corrosion resistance test as that of Example 1 was conducted. As a result of the test, neither corrosion nor breakage of the thin stainless steel film was observed.

EXAMPLE 7

ABS resin (of a plating grade) was used as a base material (member 4). The base material was a mold material, having a smooth surface.

To obtain a reflective surface of as high-degree as an electro plating, the surface of the base material was cleaned and degraded using alcohol. Then, a polyester-based resin film 1 was formed on the base material by spraying to a thickness of $15\text{--}25\mu\text{m}$. The resin film was baked at 75°C for one hour so that the base material was not deformed.

Then, a thin metal film 2 was formed by DC-magnetron-sputtering using a target of SUS304 to a thickness of about $0.08\mu\text{m}$. The target electric current

was 3A, and the vacuum pressure was 1.5×10^{-3} Torr. The film formation time period was 40 seconds.

Then, a clear acrylic resin paint was coated by spraying on the thin metal film 2 to a thickness of $20 \mu\text{m}$ and was baked at 70°C for one hour to obtain a clear protective film 3.

A visual test, a warm water immersion test and a corrosion resistance test similar to those described in Example 1 were conducted, and the same results as those for Example 1 were obtained.

EXAMPLE 8

The same test piece as that of Example 7 was prepared, except that the clear protective film was not formed after formation of the thin stainless steel film.

Then, the same corrosion resistance test as that of Example 1 was conducted. As a result of the test, neither corrosion nor breakage of the thin stainless steel film was observed.

COMPARATIVE EXAMPLE 1

The same test piece as that of Example 1 was prepared, except that a thin aluminum film of a thickness of $0.08 \mu\text{m}$ was formed as the thin metal film.

Then, the same corrosion resistance test as that of Example 1 was conducted. As a result of the test, the aluminum at the cross-cut portion disappeared after 200 hours.

COMPARATIVE EXAMPLE 2

The same test piece as that of Comparative Example 1 was prepared, except that a clear protective film was not formed after formation of the thin

aluminum film.

Then, the same corrosion resistance test as that of Example 1 was conducted. As a result of the test, the thin aluminum film disappeared after 168 hours.

5 From the test results of Comparative Example 1 and Comparative Example 2, it was found that the thin aluminum film could not maintain the corrosion resistance if the thin aluminum film was not coated with and protected by the protective film.

COMPARATIVE EXAMPLE 3

10 The same test piece as that of Example 7 was prepared, except that a thin aluminum film of a thickness of 0.08 μm was formed as the thin metal film.

Then, the same corrosion resistance test as that of Example 1 was conducted. As a result of the test, the aluminum at the cross-cut portion disappeared after 200 hours (the same as that of Comparative Example 1).

15 COMPARATIVE EXAMPLE 4

The same test piece as that of Comparative Example 3 was prepared, except that a clear protective film was not formed after formation of the thin aluminum film.

20 Then, the same corrosion resistance test as that of Example 1 was conducted. As a result of the test, the thin aluminum film disappeared after 168 hours (the same as that of Comparative Example 2).

EXAMPLE 9

25 The same test piece as that of Example 1 was prepared, except that the preparation conditions were changed as follows. In formation of the thin metal film, a sintered target containing 20% by weight of aluminum and 80% by weight

of titanium was used to form the thin metal film of a thickness of about $0.05\ \mu\text{m}$. The target electric current was 3A, and the film formation pressure was 2.5×10^{-3} Torr, and the coating period of time was 3 minutes.

5 A visual test, a warm water immersion test and a corrosion resistance test were conducted.

According to the visual test, the thin titanium alloy film did not have cracks and had an external appearance similar to that of electro chromium plating.

10 In the warm water immersion test, there was no change in the external appearance. Further, a tape was attached to the surface and then detached from the surface. As a result of the tape test, there was no separation of the clear protective film from the thin metal film.

15 In the corrosion resistance test, the clear protective film was cross-cut and was subjected to a salt water spray test for 1000 hours. As a result of the test, neither corrosion of the thin titanium alloy film from the cross-cut portion nor breakage of the thin titanium alloy film was observed.

EXAMPLE 10

The same test piece as that of Example 9 was prepared, except that the clear protective film was not formed after formation of the thin titanium alloy film.

20 Then, the same corrosion resistance test as that of Example 9 was conducted. As a result of the test, neither corrosion of the thin titanium alloy film from the cross-cut portion nor breakage of the thin titanium alloy film was observed.

EXAMPLE 11

The same test piece as that of Example 9 was prepared, except that in formation of the thin metal film a sintered target containing 50% by weight of aluminum and 50% by weight of titanium was used.

5 Then, the same tests as those of Example 9 were conducted. The results of the visual test, the warm water immersion test and the corrosion resistance test were the same as those of Example 9 except that the external appearance was a little brighter than that of Example 9.

EXAMPLE 12

10 The same test piece as that of Example 11 was prepared, except that the clear protective film was not formed after formation of the thin titanium alloy film.

Then, the same corrosion resistance test as that of Example 9 was conducted. As a result of the test, neither corrosion nor breakage of the thin titanium alloy film was observed.

15 EXAMPLE 13

The same test piece as that of Example 9 was prepared, except that in formation of the thin metal film a sintered target containing 80% by weight of aluminum and 20% by weight of titanium was used.

20 Then, the same tests as those of Example 9 were conducted. The results of the visual test, the warm water immersion test and the corrosion resistance test were the same as those of Example 9 except that the color was almost that of pure aluminum and that the reflectivity was high.

EXAMPLE 14

25 The same test piece as that of Example 13 was prepared, except that the clear protective film was not formed after formation of the thin titanium alloy

film.

Then, the same corrosion resistance test as that of Example 9 was conducted. As a result of the test, neither corrosion nor breakage of the thin titanium alloy film was observed.

5 EXAMPLE 15

10 The same test piece as that of Example 9 was prepared, except the following test conditions was. In formation of the thin metal film, a sintered target containing 50% by weight of aluminum and 50% by weight of titanium and a cathode arc-type ion plating apparatus were used and the thin metal film of the thickness of about $1\mu\text{m}$ was formed. The argon pressure was 30 mTorr, the cathode electric current was 50A, and the coating period of time was 1 hour. The temperature of the base material was controlled so that the thin metal film formation temperature did not exceed the baking temperature of the under resin film.

15 Then, the same tests as those of Example 9 were conducted. The results of the visual test, the warm water immersion test and the corrosion resistance test were the same as those of Example 9.

EXAMPLE 16

20 The same test piece as that of Example 15 was prepared, except that the clear protective film was not formed after formation of the thin titanium alloy film.

Then, the same corrosion resistance test as that of Example 9 was conducted. As a result of the test, neither corrosion nor breakage of the thin titanium alloy film was observed.

25 EXAMPLE 17

ABS resin (of a plating grade) was used as a base material (member 4).
The base material was a mold material, having a smooth surface.

To obtain a reflective surface of as high-degree as an electro plating, the
surface of the base material was cleaned and degraded using alcohol. Then, a
5 polyester-based resin film 1 was formed on the base material by spraying to a
thickness of 15–25 μm . The resin film was baked at 75°C for one hour so that the
base material was not deformed.

Then, a thin metal film 2 was formed by DC-magnetron-sputtering to a
thickness of about 0.05 μm using a sintered target containing 50% by weight of
10 aluminum and 50% by weight of titanium. The target electric current was 3A,
and the film formation pressure was 2.5×10^{-3} Torr. The film formation time
period was 3 minutes.

Then, a clear paint of acrylic resin was coated by spraying on the thin
metal film 2 to a thickness of 20 μm and was baked at 70 °C for one hour to
15 obtain a clear protective film 3.

A visual test, a warm water immersion test and a corrosion resistance test
like Example 9 were conducted, and the same results as those of Example 9 were
obtained.

EXAMPLE 18

20 The same test piece as that of Example 17 was prepared, except that the
clear protective film was not formed after formation of the thin titanium alloy
film.

Then, the same corrosion resistance test as that of Example 9 was
conducted. As a result of the test, neither corrosion nor breakage of the thin

titanium alloy film was observed.

EXAMPLE 19

The same test piece as that of Example 1 was prepared, except the following test conditions. A thin metal film 2 was formed by

5 DC-magnetron-sputtering using a melted target of hastelloy X containing 50% by weight of nickel, 22% by weight of chromium, 18.5% by weight of iron, and tungsten and cobalt etc. to a thickness of about $0.05 \mu\text{m}$. The target electric current was 3A, and the film formation pressure was 2.5×10^{-3} Torr, and the coating period of time was 20 seconds.

10 A visual test, a warm water immersion test and a corrosion resistance test were conducted.

According to the visual test, the thin nickel alloy film did not have cracks and had an external appearance similar to that of electro chromium plating.

15 In the warm water immersion test, there was no change in the external appearance. Further, a tape was attached to the surface and then detached from the surface. As a result of the tape test, there was no separation of the clear protective film from the thin metal film.

20 In the corrosion resistance test, the clear protective film was cross-cut and was subjected to a salt water spray test for 1000 hours. As a result of the test, neither corrosion of the thin nickel alloy film from the cross-cut portion nor breakage of the thin nickel alloy film was observed.

EXAMPLE 20

The same test piece as that of Example 19 was prepared, except that the clear protective film was not formed after formation of the thin nickel alloy film.

Then, the same corrosion resistance test as that of Example 19 was conducted. As a result of the test, neither corrosion of the thin nickel alloy film from the cross-cut portion nor breakage of the thin nickel alloy film was observed.

5 EXAMPLE 21

The same test piece as that of Example 19 was prepared, except that in formation of the thin metal film a melted target of inconel 600 containing 76% by weight of nickel, 15.5% by weight of chromium, 8.0% by weight of iron, and manganese, silicon, and carbon etc. was used.

10 Then, the same tests as those of Example 19 were conducted. The results of the visual test, the warm water immersion test and the corrosion resistance test were found to be the same as those of Example 19 except that the external appearance was a little brighter than that of Example 19.

EXAMPLE 22

15 The same test piece as that of Example 21 was prepared, except that the clear protective film was not formed after formation of the thin nickel alloy film.

Then, the same corrosion resistance test as that of Example 19 was conducted. As a result of the test, neither corrosion nor breakage of the thin nickel alloy film was observed.

20 EXAMPLE 23

The same test piece as that of Example 19 was prepared, except the following test conditions. In formation of the thin metal film, a melted target of inconel 600 containing 76% by weight of nickel, 15.5% by weight of chromium, 8.0% by weight of iron, and manganese, silicon, and carbon etc. was used and the
25 thin metal film of the thickness of about $0.5 \mu\text{m}$ was formed. The argon pressure

was 30 mTorr, the cathode electric current was 50A, and the coating period of time was 20 minutes. The temperature of the base material was controlled so that the thin metal film formation temperature did not exceed the baking temperature of the under resin film.

5 Then, the same tests as those of Example 19 were conducted. The results of the visual test, the warm water immersion test and the corrosion resistance test were found to be the same as those of Example 19.

EXAMPLE 24

10 The same test piece as that of Example 23 was prepared, except that the clear protective film was not formed after formation of the thin nickel alloy film.

 Then, the same corrosion resistance test as that of Example 19 was conducted. As a result of the test, neither corrosion nor breakage of the thin nickel alloy film was observed.

EXAMPLE 25

15 ABS resin (of a plating grade) was used as a base material (member 4). The base material was a mold material, having a smooth surface.

 To obtain a reflective surface of as high-degree as an electro plating, the surface of the base material was cleaned and degraded using alcohol. Then, a polyester-based resin film 1 was formed on the base material by spraying to a
20 thickness of 15-25 μ m. The resin film was baked at 75°C for one hour so that the base material was not deformed.

 Then, a thin metal film 2 was formed by DC-magnetron-sputtering to a thickness of about 0.05 μ m using a melted target of inconel 600 containing 76% by weight of nickel, 15.5% by weight of chromium, 8.0% by weight of iron, and
25 manganese, silicon, and carbon etc. The target electric current was 3A, and the

film formation pressure was 2.5×10^{-3} Torr. The film formation period of time was 20 seconds.

Then, a clear paint of acrylic resin was coated by spraying on the thin metal film 2 to a thickness of $20 \mu\text{m}$ and was baked at 70°C for one hour to obtain a clear protective film 3.

A visual test, a warm water immersion test and a corrosion resistance test like Example 19 were conducted, and the same results as those of Example 19 were obtained.

EXAMPLE 26

The same test piece as that of Example 25 was prepared, except that the clear protective film was not formed after formation of the thin nickel alloy film.

Then, the same corrosion resistance test as that of Example 19 was conducted. As a result of the test, neither corrosion nor breakage of the thin nickel alloy film was observed.

EXAMPLE 27

An aluminum alloy cast AC4C (Al-Si-Mg base) was used as a base material (metal member 4). This material was a cast, having surface roughness of $100\text{--}200 \mu\text{m}$.

Then, a chemical conversion coating was formed on the surface of the base material to increase corrosion resistance. Then, an electrostatic epoxy powder coating was conducted at the electric voltage condition of $40\text{--}60 \text{ kV}$ and was baked at 170°C for 60 minutes to form a resin film 1 of a thickness of about $120 \mu\text{m}$. Further, to obtain a smoother surface, a coating of a solvent based acrylic resin was formed to a thickness of about $25 \mu\text{m}$ and was baked at 145°C

for 60 minutes.

Then, a thin metal film 2' was formed by DC-magnetron-sputtering to a thickness of about $0.05 \mu\text{m}$ using a melted target of hastelloy X containing 50% by weight of nickel, 22% by weight of chromium, 18.5% by weight of iron, and tungsten and cobalt etc. The target electric current was 3A, and the film formation pressure was 2.5×10^{-3} Torr, and the coating period of time was 20 seconds.

Then, a clear colored paint of acrylic resin-silicone-based resin containing a sulphur dye (Sulphur Blue 4R) in the amount of 1% by weight and mixed at 100 rpm for 5 minutes was coated by spraying on the thin metal film 2' to a thickness of $20 \mu\text{m}$ and was baked at 80°C for 60 minutes to obtain a clear colored protective film 3'.

A visual test, a warm water immersion test and a corrosion resistance test were conducted.

According to the visual test, the thin nickel alloy film did not cause a crack, had a metallic reflection and had a blue color.

In the warm water immersion test, the test piece was immersed in warm water at 60°C for 72 hours. As a result of the test, there was no change in the external appearance. Further, a tape was attached to the surface and then detached from the surface. As a result of the tape test, there was no separation of the clear colored protective film from the thin metal film.

In the corrosion resistance test, the clear colored protective film was cross-cut and was subjected to a salt water spray test for 1000 hours. As a result of the test, neither corrosion of the thin nickel alloy film from the cross-cut

portion nor breakage of the thin nickel alloy film was observed.

EXAMPLE 28

5 The same test piece as that of Example 27 was prepared, except that in formation of the thin metal film a target containing 30% by weight of titanium and 70% by weight of aluminum was used.

Then, the same tests as those of Example 27 were conducted. The results of the visual test, the warm water immersion test and the corrosion resistance test were the same as those of Example 27.

EXAMPLE 29

10 Up to formation of the resin film 1, the same procedure as that of Example 27 was followed.

Then, a thin metal film 2' was formed by a cathode arc-type ion plating apparatus to a thickness of about $0.5 \mu\text{m}$ using a melted target of inconel 600 containing 76% by weight of nickel, 15.5% by weight of chromium, 8.0% by weight of iron, and manganese, silicon, and carbon etc. The argon pressure was 15 30 mTorr, the cathode electric current was 50A, and the coating period of time was 20 minutes. The temperature of the base material was controlled so that the thin metal film formation temperature did not exceed the baking temperature of the under resin film.

20 Then, a clear colored paint of acrylic resin-silicone-based resin containing a mordant dye (Yellow) in the amount of 2% by weight and mixed, was coated by spraying on the thin metal film 2' to a thickness of $20 \mu\text{m}$ and was baked at 80°C for 60 minutes to obtain a clear colored protective film 3'. The yellow dye had a good adhesion to the thin metal film containing chromium and iron, etc.

A visual test, a warm water immersion test and a corrosion resistance test were conducted. The similar test results as those of Example 27 were obtained. More particularly, the thin nickel alloy film did not crack and had a metallic reflection, and the nickel alloy film had a gold color due to the metallic reflection and the yellow color.

EXAMPLE 30

ABS resin (of a plating grade) was used as a base material (member 4). The base material was a mold material, having a smooth surface.

To obtain a reflective surface of as high-degree as an electro plating, the surface of the base material was cleaned and degraded using alcohol. Then, a polyester-based resin film 1 was formed on the base material by spraying to a thickness of $15\text{--}25\ \mu\text{m}$. The resin film was baked at 75°C for one hour so that the base material was not deformed.

Then, a thin metal film 2' was formed by DC-magnetron-sputtering to a thickness of about $0.05\ \mu\text{m}$ using a melted target of inconel 600 containing 76% by weight of nickel, 15.5% by weight of chromium, 8.0% by weight of iron, and manganese, silicon, and carbon etc. The target electric current was 3A, and the film formation pressure was 2.5×10^{-3} Torr, and the coating period of time was 20 seconds.

Then, a clear colored paint of acrylic resin-silicone-based resin containing a sulphur dye (Sulphur Yellow GG) in the amount of 2% by weight and mixed for 5 minutes, was coated by spraying on the thin metal film 2' to a thickness of $20\ \mu\text{m}$ and was baked at 80°C for 60 minutes to obtain a clear colored protective film 3'.

A visual test, a warm water immersion test and a corrosion resistance test were conducted. The test results were the same as those of Example 29.

EXAMPLE 31

Up to formation of the resin film 1, the same procedure as that of Example 27 was followed.

Then, a thin metal (aluminum) film 2' was formed by DC-magnetron-sputtering to a thickness of about 0.08 μ m using an aluminum melted target.

Then, a clear colored protective film 3' was formed in the same way as Example 29.

A visual test and a warm water immersion test were conducted. Test results were the same as those of Example 29.

EXAMPLE 32

Up to formation of the thin metal film 2', the same procedure as that of Example 31 was followed.

Then, a clear colored paint of acryl-silicone-based resin containing yellow inorganic pigment of lead chromate in the amount of 5% by weight and mixed at 200 rpm, was coated by spraying on the thin metal film 2' to a thickness of 20 μ m and was baked at 80 °C for 60 minutes to obtain a clear colored protective film 3'.

A visual test and a warm water immersion test were conducted. The similar test results as those of Example 31 were obtained. More particularly, the thin aluminum film did not cause a crack, had a metallic reflection and had a gold color due to the metallic reflection and the yellow color. Further, a grain pattern was seen which offered a design different from that of Example 31.

According to the present invention, the following technical advantages are obtained:

5 The bright surface structure according to the present invention and the structure manufactured by the method according to the present invention have a good ductility suppressing a crack generation and a good corrosion resistance.

Further, since the structure is manufactured by a dry plating method, the structure and the method therefor do not require any waste water treatment.

10 Although the present invention has been described with reference to specific exemplary embodiments, it will be appreciated by those skilled in the art that various modifications and alterations can be made to the particular embodiments shown without materially departing from the novel teachings and advantages of the present invention. Accordingly, it is to be understood that all such modifications and alterations are included within the spirit and scope of the present invention as defined by the following claims.

WHAT IS CLAIMED IS:

1 1. A bright surface structure formed on a member made from a metal or a
2 resin, said structure comprising:

3 (a) a resin film coated on said member; and

4 (b) a thin metal film formed on said resin film, said thin metal film being
5 made from a material selected from the group consisting of stainless steel, a
6 titanium alloy and a nickel alloy, wherein said metal film has a smooth surface.

1 2. A structure according to claim 1, further comprising:

2 (c) a clear protective film coated on said thin metal film.

1 3. A bright surface structure formed on a member made from a metal or a
2 resin, said structure comprising:

3 (a) a resin film coated on said member;

4 (b') a thin metal film formed on said resin film, said thin metal film being
5 made from a material selected from the group consisting of stainless steel, a
6 titanium alloy, a nickel alloy, aluminum, titanium and chromium, wherein said
7 metal film has a smooth surface, and

8 (c') a clear colored protective film coated on said thin metal film.

1 4. A structure according to any one of claims 1, 2 and 3, wherein said
2 material is austenitic stainless steel.

1 5. A structure according to any one of claims 1, 2 and 3, wherein said

2 material is a titanium alloy containing 20–80% by weight of titanium.

1 6. A structure according to any one of claims 1, 2 and 3, wherein said
2 material is a titanium alloy containing 20–80% by weight of aluminum.

1 7. A structure according to any one of claims 1, 2 and 3, wherein said
2 material is a nickel alloy containing 30–80% by weight of nickel.

1 8. A structure according to any one of claims 1, 2 and 3, wherein said
2 material is a nickel alloy containing 15–25% by weight of chromium.

1 9. A structure according to any one of claims 1, 2 and 3, wherein said
2 thin metal film is made from stainless steel or a titanium alloy and has a thickness
3 of 0.03–1.0 μm .

1 10. A structure according to any one of claims 1, 2 and 3, wherein said
2 thin metal film is made from nickel alloy and has a thickness of 0.03–0.5 μm .

1 11. A structure according to claim 2, wherein said clear protective film
2 has a thickness of 5–20 μm .

1 12. A structure according to claim 3, wherein said clear colored protective
2 film has a thickness of 20–40 μm .

1 13. A structure according to claim 3, wherein said clear colored protective

2 film is made from clear resin comprising a pigment or a dye.

1 14. A structure according to claim 13, wherein said clear resin is selected
2 from acryl-based, urethan-based or epoxy-based resins.

1 15. A structure according to claim 13, wherein said pigment is selected
2 from carbon-based, lead chromate-based, iron(II) ferrocyanide-based,
3 cobalt-based, or chromium oxide-based pigments.

1 16. A structure according to claim 13, wherein said pigment is selected
2 from thren-based, quinacrine staining-based, isoindolinone-based, or metal
3 complex pigments.

1 17. A structure according to claim 13, wherein said dye is selected from
2 an acid dye, a mordant dye, a basic dye, a disperse dye, an edible dye, a direct
3 dye or a sulphur dye.

1 18. A structure according to claim 1, wherein said member made from a
2 metal is an aluminum wheel for an automobile.

1 19. A structure according to claim 1, wherein said member made from a
2 resin is a front grille, a garnish, or an emblem of an automobile.

1 20. A method for manufacturing a bright surface structure formed on a
2 member made from a metal or a resin, said method comprising:

3 (a) coating a resin film on said member such that said resin film has a
4 smooth surface;

5 (b) forming a thin metal film on said resin film, said thin metal film being
6 made from a material selected from the group consisting of stainless steel, a
7 titanium alloy and a nickel alloy, wherein said metal film has a smooth surface.

1 21. A method according to claim 20, further comprising:

2 (c) coating a clear protective film on said metal film.

1 22. A method for manufacturing a bright surface structure formed on a
2 member made from a metal or a resin, said method comprising:

3 (a) coating a resin film on said member such that said resin film has a
4 smooth surface;

5 (b') forming a thin metal film on said resin film, said thin metal film being
6 made from material selected from the group consisting of stainless steel, a
7 titanium alloy and a nickel alloy, aluminum, titanium and chromium, wherein said
8 metal film has a smooth surface; and

9 (c') coating a clear colored protective film on said thin metal film.

1 23. A method according to any one of claims 20, 21 and 22, wherein said
2 resin film is coated by powder coating.

1 24. A method according to any one of claims 20, 21 and 22, wherein said
2 material is austenitic stainless steel.

1 25. A method according to any one of claims 20, 21 and 22, wherein said
2 material is a titanium alloy containing 20-80% by weight of titanium.

1 26. A method according to any one of claims 20, 21 and 22, wherein said
2 material is a titanium alloy containing 20-80% by weight of aluminum.

1 27. A method according to any one of claims 20, 21 and 22, wherein said
2 material is a nickel alloy containing 30-80% by weight of nickel.

1 28. A method according to any one of claims 20, 21 and 22, wherein said
2 material is a nickel alloy containing 15-25% by weight of chromium.

1 29. A method according to any one of claims 20, 21 and 22, wherein said
2 thin metal film is formed by any one of cathode arc-type ion plating and
3 sputtering.

1 30. A method according to any one of claims 20, 21 and 22, wherein said
2 thin metal film is made from stainless steel or a titanium alloy and has a thickness
3 of 0.03-1.0 μ m.

1 31. A method according to any one of claims 20, 21 and 22, wherein said
2 thin metal film is made from a nickel alloy and has a thickness of 0.03-0.5 μ m.

1 32. A method according to claim 21, wherein said clear protective film
2 has a thickness of 5-20 μ m.

1 33. A method according to claim 22, wherein said clear colored protective
2 film has a thickness of 20-40 μ m.

1 34. A method according to claim 22, wherein said clear colored protective
2 film is made from a clear resin comprising a pigment or a dye.

1 35. A method according to claim 34, wherein said clear resin is selected
2 from acryl-based, urethan-based or epoxy-based resins.

1 36. A method according to claim 34, wherein said pigment is selected
2 from carbon-based, lead chromate-based, iron(II) ferrocyanide-based,
3 cobalt-based, or chromium oxide-based pigments.

1 37. A method according to claim 34, wherein said pigment is selected
2 from thren-based, quinacrine staining-based, isoindolinone-based, or metal
3 complex pigments.

1 38. A method according to claim 34, wherein said dye is selected from
2 an acid dye, a mordant dye, a basic dye, a disperse dye, an edible dye, a direct
3 dye or a sulphur dye.

ABSTRACT OF THE DISCLOSURE

A bright surface structure formed on a member made from metal or resin.

A first structure includes: (a) a resin film coated on the member, and (b) a thin metal film formed on the resin film. The thin metal film is made from any one of stainless steel, titanium alloy and nickel alloy. A second structure further includes (c) a clear protective film coated on the thin metal film. A third structure includes: (b') a thin metal film formed on the resin film of (a), and (c') a clear colored protective film coated on the thin metal film of (b'). The thin metal film of (b') is made from any one of stainless steel, titanium alloy, nickel alloy, aluminum, titanium and chromium.

FIG. 1

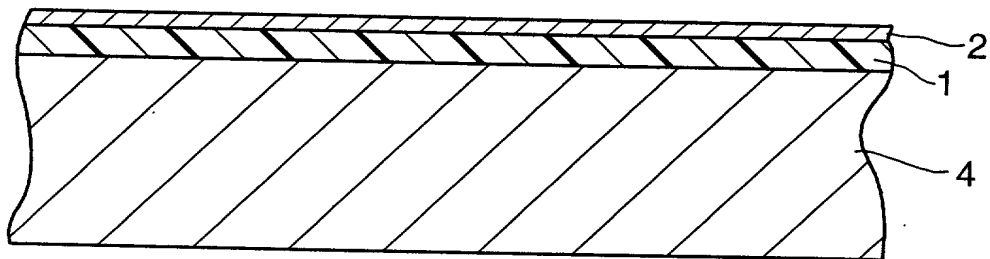


FIG. 2

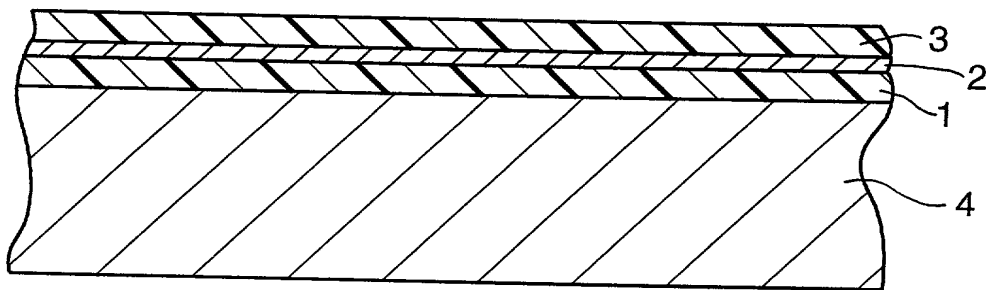
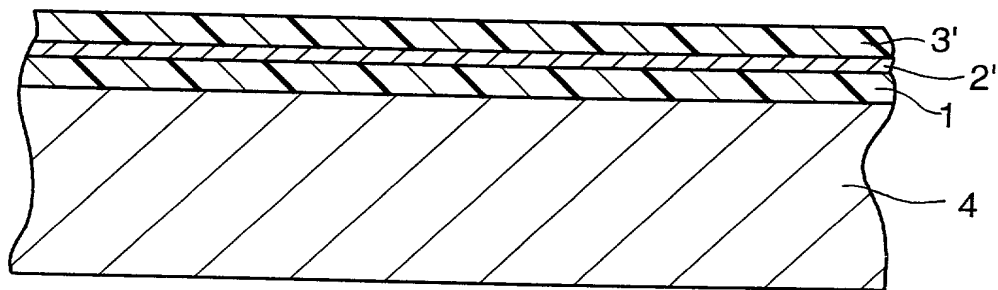


FIG. 3



FOR UTILITY/DESIGN
CIP/PCT NATIONAL/PLANT
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DECLARATIONS

RULE 63 (37 C.F.R. 1.63)
DECLARATION AND POWER OF ATTORNEY
FOR PATENT APPLICATION
IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

PM & S
FORM

As a below named inventor, I hereby declare that my residence, post office address and citizenship are as stated below next to my name, and I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the INVENTION ENTITLED BRIGHT SURFACE
STRUCTURE AND A MANUFACTURING METHOD THEREOF

the specification of which (CHECK applicable BOX(ES))
X ☒ is attached hereto.
BOX(ES) ☐ was filed on _____ as U.S. Application No. _____ /
☐ was filed as PCT International Application No. PCT/ _____ / on _____

and (if applicable to U.S. or PCT application) was amended on _____
I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above. I acknowledge the duty to disclose all information known to me to be material to patentability as defined in 37 C.F.R. 1.56 I hereby claim foreign priority benefits under 35 U.S.C. 119/365 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate filed by me or my assignee disclosing the subject matter claimed in this application and having a filing date (1) before that of the application on which priority is claimed, or (2) if no priority claimed, before the filing date of this application:

PRIOR FOREIGN APPLICATION(S)			Date first Laid- open or Published	Date Patented or Granted	Priority Claimed	
Number	Country	Day/MONTH/Year Filed			Yes	No
H11-264172	Japan	17/09/1999			X	
H11-270072	Japan	24/09/1999			X	

I hereby claim domestic priority benefit under 35 U.S.C. 119/120/365 of the indicated United States applications listed below and PCT international applications listed above or below and, if this is a continuation-in-part (CIP) application, insofar as the subject matter disclosed and claimed in this application is in addition to that disclosed in such prior applications, I acknowledge the duty to disclose all information known to me to be material to patentability as defined in 37 C.F.R. 1.56 which became available between the filing date of each such prior application and the national or PCT international filing date of this application:

PRIOR U.S. PROVISIONAL, NONPROVISIONAL AND/OR PCT APPLICATION(S)		Status	Priority Claimed	
Application No. (series code/serial no.)	Day/MONTH/Year Filed	pending, abandoned, patented	Yes	No

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

And I hereby appoint Pillsbury Madison & Sutro LLP, Intellectual Property Group, 1100 New York Avenue, N.W., Ninth Floor, East Tower, Washington, D.C. 20005-3918, telephone number (202) 861-3000 (to whom all communications are to be directed), and the below-named persons (of the same address) individually and collectively my attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith and with the resulting patent, and I hereby authorize them to delete names/numbers below of persons no longer with their firm and to act and rely on instructions from and communicate directly with the person/assignee/attorney/firm/ organization who/which first sends/sent this case to them and by whom/which I hereby declare that I have consented after full disclosure to be represented unless/until I instruct the above firm and/or a below attorney in writing to the contrary.

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First		Middle Initial		Family Name	
Naoaki		KITAGAWA			
Residence	Zushi-shi, Kanagawa-ken		Japan		Japan
	City		State/Foreign Country		Country of Citizenship
Post Office Address	7-11-48, Kotsubo, Zushi-shi, Kanagawa-ken, 249-0008 Japan				
(include Zip Code)					

(2) INVENTOR'S SIGNATURE: Shinichi Okabe Date: September 6, 2000

First		Middle Initial		Family Name	
Shinichi		OKABE			
Residence	Yamato-shi, Kanagawa-ken		Japan		Japan
	City		State/Foreign Country		Country of Citizenship
Post Office Address	4-10-9, Minamirinkan, Yamato-shi, Kanagawa-ken, 242-0006 Japan				
(include Zip Code)					

(FOR ADDITIONAL INVENTORS, check box ☐ to attach PAT 116-2 same information for each re signature, name, date, citizenship, residence and address.)